

An Information Service of the Division of Medical Assistance

North Carolina Medicaid Pharmacy

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Policy and Procedures for Prescribing Synagis for RSV Season 2005-2006

For the upcoming RSV season, Synagis will not require prior approval (PA) for Medicaid recipients. However, the responsibility for appropriate usage of Synagis will be placed on prescribers and pharmacy providers. The clinical criteria utilized in this policy are consistent with currently published American Academy of Pediatrics guidelines (http://aappolicy.aappublications.org/cgi/content/full/pediatrics; 112/6/1442). Please ensure that the person completing the Synagis criteria form has verified that the conditions exist and are accurate. If a patient does not fit the published criteria and you still wish to prescribe Synagis, you must submit your request to DMA on the *Request for Medical Review for Synagis Outside of Criteria* form and fax the request to DMA at 919-657-8843.

The start of the Synagis season is October 15, 2005. No more than 5 monthly doses of Synagis can be obtained by using these forms. The number of doses should be adjusted if an infant received the first dose prior to a hospital discharge. Delays in getting a request processed can occur if the patient does not have a Medicaid identification number or the form is not complete.

The criteria form must be signed by the prescriber and submitted to the pharmacy distributor of choice. The pharmacy distributor must mail a copy of the submitted form to DMA. Please mail to:

N.C. Division of Medical Assistance Pharmacy Program 1985 Umstead Drive 2501 Mail Service Center Raleigh, N.C. 27699

The Request for Medical Review for Synagis Outside of Criteria form must be signed by the prescriber and faxed to DMA at 919-657-8843.

Please refer to the following guidelines when submitting a request:

• For the following four diagnoses, DOB must be on or after 10/15/03:

Chronic Lung Disease (Bronchopulmonary Dysplasia)

The infant has Chronic Lung Disease (bronchopulmonary dysplasia) and has necessitated treatment (supplemental oxygen, bronchodilator, diuretic, corticosteroid) in the six months before the start of the season.

<u>Hemodynamically Significant Congenital Heart Disease</u>

Infants less than 12 months of age who are most likely to benefit include those receiving medication to control CHF, moderate to severe pulmonary hypertension, and/or cyanotic heart disease.

Infants NOT at increased risk from RSV who generally should NOT receive immunoprophylaxis include: hemodynamically insignificant heart disease such as secundum atrial/septal defect, small VSD, pulmonic stenosis, uncomplicated

aortic stenosis, mild coarctation of the aorta, PDA, lesions adequately corrected by surgery unless the infant continues on medication for CHF, mild cardiomyopathy where the infant is not receiving medical therapy.

Cystic Fibrosis

The infant has Cystic Fibrosis and either requires chronic oxygen or has been diagnosed with nutritional failure.

Severe Congenital Immunodeficiency

Severe combined immunodeficiency disease or severe acquired immunodeficiency syndrome.

Infant is born at an EGA of:

 \leq 28 weeks and DOB is on or after 10/15/04 29-32 weeks and DOB is on or after 4/15/05

■ If born between 32 weeks and 1 day and 35 weeks and 0 days gestation, must be less than 6 months of age (DOB on or after 4/15/05) at the start of the season and have two or more defined risk factors:

☐ School-age Siblings
☐ Attends Day Care
☐ Severe Neuromuscular Disease
☐ Exposure to prolonged wood burning heaters which are the primary source of
heat for the family. Tobacco smoke is NOT a risk factor because it can be
controlled by the family.
☐ Congenital abnormalities of the airways.

Request for Medical Review for Synagis Outside of Criteria

This form will be used for patients who do not explicitly meet the guidelines whose providers still wish to prescribe Synagis. Please fill out the requested information and fax to DMA at 919-657-8843. <u>PLEASE NOTE THAT THIS</u> IS THE ONLY FORM THAT PRESCRIBERS SHOULD FAX TO DMA.

Medicaid will allow Synagis claims processing to begin on October 10, 2005 to allow sufficient time for pharmacies to provide Synagis by the start of the season on October 15, 2005. Payment of Synagis claims prior to October 10, 2005 and after March 15, 2006 will not be allowed. Physicians and pharmacy providers are subject to audits of Synagis records by DMA Program Integrity.

The Synagis Criteria Form and the Request for Medical Review for Synagis Outside of Criteria Form are available on the DMA website at http://www.dhhs.state.nc.us/dma/prov.htm

Family Planning Waiver

On October 1, 2005, the N.C. Medicaid program will implement the new Family Planning Waiver program. This is a Medicaid program designed to reduce unintended pregnancies and improve the wellbeing of children and families in North Carolina by extending eligibility for family planning services to eligible women between the ages of 19 through 55 and men ages 19 through 60 whose income is at or below 185% of the federal poverty level.

Objectives of the Family Planning Waiver are:

- Increase the number of reproductive age women and men receiving either Title XIX or Title X funded family planning services by improving access to and use of Medicaid family planning services.
- Reduce the number of inadequately spaced pregnancies by women in the target group thus improving birth outcomes and health of these women.
- Reduce the number of unintended and unwanted pregnancies among women eligible for Medicaid.
- Impact positively the utilization of and "continuation rates" for contraceptive use among the target population.
- Increase the use of more effective methods of contraception in the target population.

Recipients have the freedom of choice in deciding to receive or reject any family planning service.

Eligible Recipients

A new Medicaid eligibility category, MAF-D, has been created for the waiver. The eligible recipient will be identified by a blue Medicaid card with the following statement "FAMILY PLANNING WAIVER: RECIPIENT ELIGIBLE FOR LIMITED FAMILY PLANNING SERVICES ONLY" (Sample card). Only one name will be listed per Medicaid card. Recipients eligible to receive waiver services are not eligible for Medicaid under any other current program.

Eligible recipients are:

- Women age 19 through 55 or men age 19 through 60
- Income at or below 185% of the federal poverty level
- U.S. citizens or qualified alien
- Residents of North Carolina
- Not incarcerated
- Not pregnant
- Not permanently sterilized

Patients who are eligible will be identified by a blue Medicaid card with the following statement "Family Planning Waiver" (See sample below).

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		RECIP	ENT ID		ELIOIBLI	ES FOR MEDI	CAID		INS NO	BIRTHDATE	SEX
P.O. Box 111 Any City, NC		3-45	-6789K	Ja	ane Recipie	пt			l	08-02-1971	F
Zip=12345				*** Family Planning Waiver *** Recipient Eligible For Limited Family Planning Services Only							
CASEID 10847667 CASEHEAD Jane Recipient		INS NAME POLICY NUMBER TYPE SEP 200 Caschea			EP 2005 Aschead N	lame	10847667 10	I			
Eligible Members		456 That Str That City, N									
Jane Recipient		RRECIPIENT (Signature)				(Not valid unless	signed)				
123-45-6789K	iL						`				

- There is no co-payment for services received under the Family Planning Waiver program.
- The six prescriptions limit will apply, no overrides allowed.
- All claims will be paid based on Medicaid reimbursement policy.
- Dispensing fees are based on Medicaid policy.
- Providers are not allowed to indicate brand "medically necessary" (DAW=1) for drugs where a generic is available.
- All claims must be submitted via point of sale (POS) and must have one of the approved ICD-9 codes.
- If the recipient receives several antibiotics for different STI's, all claims must be filled on the same date of service and different ICD-9 codes must be indicated.
- Contraceptives do not require an ICD-9 code.
- Compounds are not covered.

NOTE:

Providers are required to have a FP diagnosis on all prescriptions (excluding contraceptives).

Medication List

All contraceptives are eligible for the Family Planning Waiver and are identified by the family planning indicator on the drug file. In addition, the drugs and strengths identified below are covered if prescribed for one of the STI diagnoses indicated or for postoperative medication. Prescribers are not allowed to indicate brand "medically necessary" (DAW = 1) for drugs where a generic is available.

Birth control pills may be dispensed through a pharmacy. A recipient may receive up to a 3-month supply.

STI MEDICATION LIST

Medications for the Family Planning Waiver program will only be provided by prescription through the pharmacy drug program.

STI DIAGNOSIS	ICD-9 CODE	REIMBURSED ANTIBIOTICS
HERPES	CODE	Acyclovir 200mg, 400mg, 800 mg
Genital herpes	54.10	Famciclovir 125mg, 250mg, 500mg
Herpetic vulvovaginitis	54.11	Valacyclovir 500mg, 1.0gm
Herpetic ulceration of vulva	54.12	
Herpetic infection of penis	54.13	
Other	54.19	
CHLAMYDIA		Azithromycin, 250mg, 500mg, 1gm
Other specified diseases due to	78.88	Doxycycline 100mg
Chlamydia		Erythromycin 250mg, 400mg, 500mg,
Other venereal disease, Chlamydia	99.41	800mg
trachomatis		Ofloxacin 200mg, 300mg, 400mg
Other venereal disease, Lower GU site	99.53	Levofloxacin 500mg
		Tetracycline 250mg
SYPHILIS		Azithromycin 1gm
Genital syphilis (primary)	91.0	Benzathine penicillin G 2.4 million units
Primary anal syphilis	91.1	Ceftriazone 250mg
Other primary syphilis	91.2	Ciprofloxacin 500mg
Early syphilis, latent, serological	92.0	Doxycycline 100mg
relapse after treatment		Erythromycin 500mg
Early syphilis, latent, unspecified	92.9	Tetracycline 500mg
GONORRHEA		Azithromycin 250mg, 500mg, 1gm
Acute, of lower GU tract	98.0	Cefixime 400mg
Gonococcal infection (acute) of upper	98.10	Ceftriaxone 125 mg, 250mg, 500mg
GU tract, site unspecified		Ceftizoxime 500mg
Gonococcal cystitis (acute)	98.11	Cefotaxime 500mg
Gonococcal prostatitis (acute)	98.12	Cefoxitin 2gm with probenecid 1gm
Gonococcal epididymo-orchitis (acute)	98.13	Ciprofloxacin 250mg, 500mg
Gonococcal seminal vesiculitis (acute)	98.14	Cefpodoxime 200 mg
Gonococcal cervicitis (acute)	98.15	Gatifloxacin 400mg
Gonococcal endometritis (acute)	98.16	Levofloxacin 250mg
Gonococcal salpingitis, acute	98.17	Lomefloxacin 400mg

Other	98.19	Norfloxaxin 800mg
Chronic, of lower GU tract	98.2	Ofloxacin 400mg
Chronic, gonococcal infection of upper	98.30	Spectinomycin 2gm
GU tract, site unspecified	70.00	Sulfamethoxazole/TMP
Gonococcal cystitis, chronic	98.31	
Gonococcal prostatitis, chronic	98.32	
Gonococcal epididymo-orchitis, chronic	98.33	
Gonococcal seminal vesiculitis, chronic	98.34	
Gonococcal cervicitis, chronic	98.35	
Gonococcal endometritis, chronic	98.36	
Gonococcal salpingitis (chronic)	98.37	
Other	98.39	
Gonococcal arthritis	98.50	
Gonococcal synovitis and tenosynovitis	98.51	
Gonococcal bursitis	98.52	
Gonococcal spondylitis	98.53	
Other	98.59	
Gonococcal infection of pharynx	98.6	
Gonococcal infection of anus and	98.7	
rectum		
OTHER VENEREAL DISEASE		Azithromycin 250mg, 500mg, 1gm
Non-gonococcal urethritis, unspecified	99.40	Doxycycline 100mg
		Erythromycin 500mg, 800mg
		Gatifloxacin 400mg
		Levofloxacin 250mg, 500mg
		Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg
CANDIDIASIS		Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream
Of vulva and vagina	112.1	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository
	112.1 112.2	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository
Of vulva and vagina Of other urogenital sites		Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8%
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS	112.2	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified	112.2	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8%
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis	112.2 131.00 131.01	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis	112.2 131.00 131.01 131.02	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis Trichomonal prostatitis	112.2 131.00 131.01 131.02 131.03	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis Trichomonal prostatitis Other	112.2 131.00 131.01 131.02 131.03 131.09	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis Trichomonal prostatitis Other Other specified sites	131.00 131.01 131.02 131.03 131.09 131.8	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis Trichomonal prostatitis Other Other specified sites Trichomoniasis, unspecified	112.2 131.00 131.01 131.02 131.03 131.09	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm Tinidazole 2000mg
Of vulva and vagina Of other urogenital sites TRICHOMONIASIS Urogenital trichomoniasis, unspecified Trochomonal vulvovaginitis Trichomonal urethritis Trichomonal prostatitis Other Other specified sites	131.00 131.01 131.02 131.03 131.09 131.8	Levofloxacin 250mg, 500mg Ofloxacin 200mg, 300mg, 400mg Butoconazole 2% cream Miconazole 200mg suppository Terconazole 80mg suppository Terconazole cream 0.4%, 0.8% Metronidazole 250mg, 500mg, 750mg, 2gm

NOTE:
For additional information regarding STI infections and diagnosis, refer to the Center for Disease Control (CDC) Sexually Transmitted Diseases Treatment Guidelines.

POSTOPERATIVE STERILIZATION MEDICATION LIST

Medications for the Family Planning Waiver program will only be provided by prescription through the pharmacy drug program. All prescriptions for postoperative sterilization medications must include a V25.2 diagnosis.

STERILIZATION PROCEDURE	REIMBURSED ANTIBIOTICS
VASECTOMY	Amox TR-K CLV 500-125mg, 1000-
Vasectomy, unilateral or bilateral (including	62.5
postop semen examination(s))	Amoxicillin 250mg, 500mg
Ligation of vas deferens, unilateral or	Cephalexin 250mg, 500mg
bilateral	Ciprofloxacin HCL 250mg, 500mg
TUBAL LIGATION	Doxycycline 100mg
Ligation or transaction fallopian tubes	Erythromycin ES 400mg
abdominal or vaginal approach, unilateral or	Levofloxacin 500mg
bilateral	Metronidazole 500mg
Occlusion of fallopian tube(s) by device	Penicillin VK 500mg
vaginal or suprapubic approach	Sulfamethoxazole/TMP DS
Laparoscopy surgical; with fulguration of	Azithromax 250mg
oviducts (with or without transaction)	
Tubal ligation by laparoscopic surgery with	
occlusion of device (band, clip or Falope	
ring)	
STERILIZATION PROCEDURE	REIMBURSED ANALGESICS
VASECTOMY	Acetaminophen/Cod #2, #3
Vasectomy, unilateral or bilateral (including	Hydrocodone/Apap 2.5/500, 5/325,
postop semen examination(s))	5/500, 7.5/325, 7.5/500, 7.5/650,
Ligation of vas deferens, unilateral or	7.5/750, 10/325, 10/500, 10/650,
bilateral	10/660, 10/750
TUBAL LIGATION	Ibuprofen 400mg, 600mg, 800mg
Ligation or transaction fallopian tubes	Ketorolac 10mg
abdominal or vaginal approach, unilateral or	Naproxen 500mg
bilateral	Naproxen Sodium 550mg
Occlusion of fallopian tube(s) by device	Oxycodone 5mg
vaginal or suprapubic approach	Oxycodone w/Apap 2.5/325, 5/325,
Laparoscopy surgical; with fulguration of	5/325, 7.5/325, 7.5/500, 10/325, 10/650
oviducts (with or without transaction)	Propoxy-N/Apap 65/650, 100-650
Tubal ligation by laparoscopic surgery with	
occlusion of device (band, clip or Falope	
ring)	

STERILIZATION PROCEDURE	REIMBURSED ANALGESICS
VASECTOMY	Promethazine 25mg
Vasectomy, unilateral or bilateral (including	
postop semen examination(s))	
Ligation of vas deferens, unilateral or	
bilateral	
TUBAL LIGATION	
Ligation or transaction fallopian tubes	
abdominal or vaginal approach, unilateral or	
bilateral	
Occlusion of fallopian tube(s) by device	
vaginal or suprapubic approach	
Laparoscopy surgical; with fulguration of	
oviducts (with or without transaction)	
Tubal ligation by laparoscopic surgery with	
occlusion of device (band, clip or Falope	
ring)	

Medicare Part D Provider Marketing Guidelines Now Available

The final version of the Medicare Part D Marketing Guidelines is now available from the Centers for Medicare and Medicaid Services and can be found at http://www.cms.hhs.gov/pdps/. The Guidelines include information on what pharmacists can and cannot do when assisting beneficiaries with prescription drug plan (PDP) selection.

Pharmacy providers contracted with Plans and their subcontractors can:

- Provide the names of Plans with which they contract and/or participate.
- Provide information and assistance in applying for the limited income subsidy.
- Provide objective information on specific Plan formularies, based on a particular patient's medications and health care needs.
- Provide objective information regarding specific Plans, such as covered benefits, cost sharing, and utilization management tools.
- Distribute PDP marketing materials, including enrollment application forms. Note: Providers must inform individuals where they can obtain information on all available options within the service area (i.e., 1-800-MEDICARE or http://www.medicare.gov/).
- Distribute Medicare Advantage (MA) and/or Medicare Advantage-Prescription Drug (MA-PD) marketing materials, excluding enrollment application forms.

Note: Providers must inform individuals where they can obtain information on all available options within the service area (i.e., 1-800-MEDICARE or http://www.medicare.gov/).

- Refer their patients to other sources of information, such as the State Health Insurance Assistance Programs, Plan marketing representatives, CMS's website at http://www.medicare.gov/, or calling 1-800-MEDICARE.
- Print out and share information with patients from CMS's website.
- Use comparative marketing materials comparing plan information created by a non-benefit/service providing third party (See Section 10 of the Guidelines under Marketing of Multiple Lines of Business, Non-Benefit/Service-Providing Third Party Marketing Materials).

Pharmacy providers contracted with Plans and their subcontractors **cannot**:

- Direct, urge, or attempt to persuade, any prospective enrollee to enroll in a particular Plan or to insure with a particular company based on financial or any other interest of the provider (subcontractor).
- Collect enrollment applications.
- Offer inducements to persuade beneficiaries to enroll in a particular plan or organization.
- Health screen when distributing information to patients, as health screening is a prohibited marketing activity.
- Offer anything of value to induce Plan enrollees to select them as their provider.
- Expect compensation in consideration for the enrollment of a beneficiary.
- Expect compensation directly or indirectly from the Plan for beneficiary enrollment activities.
- The Guidelines are provided to help pharmacists assist beneficiaries with prescription drug plan selection and to also ensure that plan selection is in the best interest of the beneficiary.

Change in Pharmacy Co-Pay

Effective November 1, 2005, the co-pay for Medicaid prescriptions will change to \$3.00 for generic drugs. Brand name drugs will remain \$3.00. All co-pay rules will remain the same.

Providing Compounded Prescriptions for Medicaid Recipients

Medicaid pharmacy providers providing compounded prescriptions to the general public may not exclude Medicaid recipients from receiving compounded prescriptions from their pharmacies. Medicaid pharmacy providers that are the pharmacy of record must take the necessary steps to ensure that Medicaid recipients receive prescribed medications within the month of record including prescribed medications that require compounding. If the ingredients required for the compounded prescription are not available, the pharmacy of record must assist the Medicaid recipient with locating a pharmacy that has the ingredients necessary to fill the prescribed compounded prescription.

Changes in Drug Rebate Manufacturers

Additions

The following labeler has entered into a Drug Rebate Agreement and joined the rebate program effective on the date indicated below:

Code	Manufacturer	Date
15020	GSP Company	September 9, 2005

Provider Services and Enrollment Change of Address

This is a reminder that Medicaid providers must report all changes to DMA using the Medicaid Provider Change Form located at DMA's website:

http://www.dhhs.state.nc.us/dma/forms.html#prov

This includes change in address and phone, but providers now can also include a fax number and e-mail address.

The Provider Services and Enrollment Section of the Division of Medical Assistance has moved. Certified mail, UPS, or Federal Express deliveries that require a street address should be sent to the following address:

DMA Provider Services Attn: *First Name Last Name* 801 Ruggles Drive Raleigh, NC 27603 The mailing address and telephone numbers will remain the same:

DMA Provider Services Attn: *First Name Last Name* 2501 Mail Service Center Raleigh, NC 27600-2501 919-855-4050

NCLeads Update

Information related to the implementation of the new Medicaid Management Information System, *NCLeads*, scheduled for implementation in mid-2006 can be found online at http://ncleads.dhhs.state.nc.us. Please refer to this website for information, updates, and contact information related to the *NCLeads* system.

Checkwrite Schedule

September 07, 2005	October 11, 2005	November 08, 2005
September 13, 2005	October 18, 2005	November 15, 2005
September 20, 2005	October 27, 2005	November 23, 2005
September 29, 2005		

Electronic Cut-Off Schedule

September 02, 2005	October 07, 2005	November 04, 2005
September 09, 2005	October 14, 2005	November 11, 2005
September 16, 2005	October 21, 2005	November 18, 2005
September 23, 2005		

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